

The genetic architecture of target-site resistance to pyrethroid insecticides in the African malaria vectors *Anopheles gambiae* and *Anopheles coluzzii*

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Abstract

Resistance to pyrethroid insecticides is a major concern for malaria vector control, because these are the compounds used in almost all insecticide-treated bed-nets (ITNs), and are also widely used for indoor residual spraying (IRS). Pyrethroids target the voltage-gated sodium channel (VGSC), an essential component of the mosquito nervous system, but substitutions in the amino acid sequence can disrupt the activity of these insecticides, inducing a resistance phenotype. Here we use Illumina whole-genome sequence data from phase 2 of the *Anopheles gambiae* 1000 Genomes Project (Ag1000G) to provide a comprehensive account of genetic variation in the *Vgsc* gene in mosquito populations from 13 African countries. In addition to the three known *kdr*

resistance alleles, we describe 20 non-synonymous nucleotide substitutions at appreciable frequency in one or more populations that are previously unknown in *Anopheles* mosquitoes. Thirteen of these novel alleles were found to occur almost exclusively on haplotypes carrying the known L995F resistance allele (L1014F in *Musca domestica* codon numbering), and may enhance or compensate for the L995F resistance phenotype. A novel mutation I1527T, which is adjacent to a predicted pyrethroid binding site, was found in tight linkage with either of two alleles causing a V402L substitution, similar to a combination of substitutions found to cause pyrethroid resistance in several other insect species. We analyse the genetic backgrounds on which non-synonymous alleles are found, to determine which alleles have experienced recent positive selection, and to refine our understanding of the spread of resistance between species and geographical locations. We describe ten distinct *kdr* carrying haplotype groups with evidence of recent positive selection, five of which carry the known L995F resistance allele, five of which carry the known L995S resistance allele. Five of these groups are localised to a single geographical location, and five comprise haplotypes from different countries, in one case separated by over 3000 km, providing new information about the geographical distribution and spread of resistance. Two "non-*kdr*" haplotype groups with evidence of recent selection were also detected, one of which carries the novel I1527T allele, and one of which carries a novel M490I allele. We also find evidence for multiple introgression events transmitting resistance alleles between *An. gambiae* and *An. coluzzii*. We identify markers that could be used to design high-throughput, low-cost genetic assays for improved surveillance of pyrethroid resistance in the field. Our results demonstrate that the molecular basis of target-site pyrethroid resistance in malaria vectors is more complex than previously appreciated, and provide a foundation for the development of new genetic tools to track the spread insecticide resistance and improve the design of strategies for insecticide resistance management.

Introduction

Pyrethroid insecticides have been the cornerstone of malaria prevention in Africa for almost two decades [1]. Pyrethroids are currently used in all insecticide-treated bed-nets (ITNs), and are widely used in indoor residual spraying (IRS) campaigns as well as in agriculture. Pyrethroid resistance is widespread in malaria vector populations across Africa [2]. The World Health Organization (WHO) has published plans for insecticide resistance

management (IRM), which emphasise the need for improvements in both our knowledge of the molecular mechanisms of resistance and our ability to monitor them in natural populations [3].

The voltage-gated sodium channel (VGSC) is the physiological target of pyrethroid insecticides, and is integral to the insect nervous system. Pyrethroid molecules bind to sites within the protein channel and prevent normal nervous system function, causing paralysis (“knock-down”) and then death. However, amino acid substitutions at key positions within the protein alter the interaction with insecticide molecules (target-site resistance), increasing the dose of insecticide required for knock-down [4, 5]. In the African malaria vectors *Anopheles gambiae* and *An. coluzzii*, three substitutions have been found to cause pyrethroid resistance. Two of these substitutions occur in codon 995¹, with L995F prevalent in West and Central Africa [6, 7], and L995S found in Central and East Africa [8, 7]. A third substitution, N1570Y, has been found in West and Central Africa and shown to increase resistance in association with L995F [10]. However, studies in other insect species have found a variety of other *Vgsc* substitutions inducing a resistance phenotype [11, 12, 5]. To our knowledge, no studies in malaria vectors have analysed the full *Vgsc* coding sequence, thus the molecular basis of target-site resistance to pyrethroids has not been fully explored.

Basic information is also lacking about the spread of pyrethroid resistance in malaria vectors. For example, it is not clear when, where or how many times pyrethroid target-site resistance has emerged. Geographical paths of transmission, carrying resistance alleles between mosquito populations, are also not known. Previous studies have found evidence that L995F occurs on several different genetic backgrounds, suggesting multiple independent outbreaks of resistance driven by this allele [13, 14, 15, 16]. However, these studies analysed only small gene regions in a limited number of mosquito populations, and therefore had limited resolution to make inferences about relationships between haplotypes carrying this allele. It has also been shown that the L995F allele spread from *An. gambiae* to *An. coluzzii* in West Africa [Weill2000, 17, 18, 19]. However, both L995F and L995S

¹Codon numbering is given here relative to transcript AGAP004707-RA as defined in the AgamP4.4 gene annotations. A mapping of codon numbers from AGAP004707-RD (gene set: AgamP4.12) to *Musca domestica*, the system in which knock-down resistance mutations were first described [9], is given in Table 1.

84 now have wide geographical distributions [7], and to our knowledge no attempts have been
85 made to infer or track the geographical spread of either allele across Africa.

86 Here we report an in-depth analysis of genetic variation in the *Vgsc* gene, using whole-
87 genome Illumina sequence data from phase 1 of the *Anopheles gambiae* 1000 Genomes
88 Project (Ag1000G) [20]. The Ag1000G phase 1 resource includes data on nucleotide vari-
89 ation in 765 wild-caught mosquitoes sampled from 8 countries, with representation of
90 West, Central, Southern and East Africa, and of both *An. gambiae* and *An. coluzzii*.
91 We investigate variation across the complete gene coding sequence, and report popula-
92 tion genetic data for both known and novel non-synonymous nucleotide substitutions. We
93 then use haplotype data from the chromosomal region spanning the *Vgsc* gene to study
94 the genetic backgrounds carrying resistance alleles, infer the geographical spread of resis-
95 tance between mosquito populations, and provide evidence for recent positive selection.
96 Finally, we explore ways in which variation data from Ag1000G can be used to design
97 high-throughput, low-cost genetic assays for surveillance of pyrethroid resistance, with
98 the capability to differentiate and track resistance outbreaks.

99 Results

100 *Vgsc* non-synonymous nucleotide variation

101 To identify variants with a potentially functional role in pyrethroid resistance, we extracted
102 single nucleotide polymorphisms (SNPs) that alter the amino acid sequence of the VGSC
103 protein from the Ag1000G phase 1 data resource. We then computed their allele frequen-
104 cies among 9 mosquito populations defined by species and country of origin. Alleles that
105 confer resistance are expected to increase in frequency under selective pressure, and we
106 filtered the list of potentially functional variant alleles to retain only those at or above 5%
107 frequency in one or more populations (Table 1). The resulting list comprises 23 variant
108 alleles, including the known L995F, L995S and N1570Y resistance alleles, and a further 20
109 alleles not previously described in anopheline mosquitoes. We reported 15 of these novel
110 alleles in our overall analysis of the Ag1000G phase 1 data resource [20], and we extend
111 the analyses here to incorporate a SNP which alters codon 1603 and two tri-allelic SNPs
112 affecting codons 402 and 490.

Table 1. Non-synonymous nucleotide variation in the voltage-gated sodium channel gene. AO=Angola; BF=Burkina Faso; GN=Guinea; CM=Cameroon; GA=Gabon; UG=Uganda; KE=Kenya; GW=Guinea-Bissau; *Ac*=*An. coluzzii*; *Ag*=*An. gambiae*. Species status of specimens from Kenya and Guinea-Bissau is uncertain [20]. All variants are at 5% frequency or above in one or more of the 9 Ag1000G phase 1 populations, with the exception of 2,400,071 G>T which is only found in the CMAg population at 0.4% frequency but is included because another mutation (2,400,071 G>A) is found at the same position causing the same amino acid substitution (M490I); and 2,431,019 T>C (F1920S) which is at 4% frequency in GAAg but also found in CMAg and linked to L995F.

Variant				Population allele frequency (%)															
Position ¹	Ag ²	Md ³	Domain ⁴	AOAc	GHAc	BFAc	CIAc	GNAc	GW	GM	CMAg	GHAg	BFAg	GNAg	GAAg	UGAg	GQAg	FRAg	KE
2,390,177 G>A	R254K	R261	IL45	0	1	0	0	0	0	0	31	0	0	0	20	0	0	0	0
2,391,228 G>C	V402L	V410	IS6	0	13	7	8	12	0	0	0	0	0	0	0	0	0	0	0
2,391,228 G>T	V402L	V410	IS6	0	5	6	0	0	0	0	0	0	0	0	0	0	0	0	0
2,399,997 G>C	D466H	-	LI/II	0	0	0	0	0	0	0	7	0	0	0	0	0	0	0	0
2,400,071 G>A	M490I	M508	LI/II	0	0	0	0	0	0	3	0	0	0	0	0	0	0	0	19
2,400,071 G>T	M490I	M508	LI/II	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
2,402,466 G>T	G531V	G549	LI/II	0	0	0	0	0	0	0	0	0	0	0	1	0	6	0	0
2,407,967 A>C	Q697P	Q724	LI/II	0	0	0	0	0	0	0	0	0	0	0	0	0	6	0	0
2,416,980 C>T	T791M	T810	IIS1	0	1	2	0	0	0	0	0	29	15	11	0	0	0	0	0
2,422,651 T>C	L995S	L1014	IIS6	0	0	0	0	0	0	0	16	0	0	0	67	100	0	0	76
2,422,652 A>T	L995F	L1014	IIS6	84	82	85	92	88	0	0	53	100	100	100	33	0	0	0	0
2,429,556 G>A	V1507I	-	IIIL56	0	0	0	0	12	0	0	0	0	0	0	0	0	0	0	0
2,429,617 T>C	I1527T	I1532	IIIS6	0	17	13	8	12	0	0	0	0	0	0	0	0	0	0	0
2,429,745 A>T	N1570Y	N1575	LIII/IV	0	0	27	0	0	0	0	6	17	21	9	0	0	0	0	0
2,429,897 A>G	E1597G	E1602	LIII/IV	0	0	0	0	0	0	0	0	0	7	6	0	0	0	0	0
2,429,915 A>C	K1603T	K1608	IVS1	0	5	5	0	0	0	0	0	0	0	0	0	0	0	0	0
2,430,424 G>T	A1746S	A1751	IVS5	0	0	0	0	0	0	0	0	29	14	10	0	0	0	0	0
2,430,817 G>A	V1853I	V1858	COOH	0	0	0	0	0	0	0	0	54	5	6	0	0	0	0	0
2,430,863 T>C	I1868T	I1873	COOH	0	0	0	0	0	0	0	0	0	26	20	0	0	0	0	0
2,430,880 C>T	P1874S	P1879	COOH	0	3	21	35	0	0	0	0	0	0	0	0	0	0	0	0
2,430,881 C>T	P1874L	P1879	COOH	0	0	7	1	25	0	0	0	0	23	48	0	0	0	0	0
2,431,061 C>T	A1934V	A1939	COOH	0	2	11	46	50	0	0	0	0	0	0	0	0	0	0	0
2,431,079 T>C	I1940T	I1945	COOH	0	12	4	0	0	0	0	7	0	0	0	0	0	0	0	0

¹ Position relative to the AgamP3 reference sequence, chromosome arm 2L. Variants marked with an asterisk (*) failed conservative variant filters applied genome-wide in the Ag1000G phase 1 AR3 callset, but appeared sound on manual inspection of read alignments.

² Codon numbering according to *Anopheles gambiae* transcript AGAP004707-RA in geneset AgamP4.4.

³ Codon numbering according to *Musca domestica* EMBL accession X96668 [9].

⁴ Location of the variant within the protein structure. Transmembrane segments are named according to domain number (in Roman numerals) followed by ‘S’ then the number of the segment; e.g., ‘IIS6’ means domain two, transmembrane segment six. Internal linkers between segments within the same domain are named according to domain (in Roman numerals) followed by ‘L’ then the numbers of the linked segments; e.g., ‘IL45’ means domain one, linker between transmembrane segments four and five. Internal linkers between domains are named ‘L’ followed by the linked domains; e.g., ‘LI/II’ means the linker between domains one and two. ‘COOH’ means the internal carboxyl tail.

113 The two known resistance alleles affecting codon 995 had the highest overall allele fre-
114 quencies within the Ag1000G phase 1 cohort (Table 1). The L995F allele was at high
115 frequency in populations of both species from West, Central and Southern Africa . The
116 L995S allele was at high frequency among *An. gambiae* populations from Central and East
117 Africa. Both alleles were present in *An. gambiae* populations sampled from Cameroon
118 and Gabon, including some individuals with a hybrid L995F/S genotype (46/275 individ-
119 uals in Cameroon, 36/56 in Gabon). In Cameroon these alleles were in Hardy Weinberg
120 equilibrium ($\chi^2 = 0.02$, $P > 0.05$), but there was an excess of heterozygotes in Gabon (χ^2
121 $= 8.96$, $P < 0.005$), suggesting a fitness advantage for mosquitoes carrying both alleles at
122 least in some circumstances.

123 The N1570Y allele was present in Guinea, Burkina Faso (both species) and Cameroon.
124 This allele has only ever been found in association with L995F [10], and has been shown
125 to substantially increase pyrethroid resistance when it occurs in combination with L995F,
126 both in association tests of phenotyped field samples [10] and experimentally [21]. To
127 study the patterns of association among non-synonymous variants, we used haplotypes
128 from the Ag1000G phase 1 resource to compute the normalised coefficient of linkage dis-
129 equilibrium (D') between all pairs of variant alleles (Figure 1). As expected, we found
130 N1570Y in almost perfect linkage with L995F, meaning that N1570Y was almost only ever
131 found on haplotypes carrying L995F. Of the 20 novel non-synonymous alleles, 13 also oc-
132 curred almost exclusively in combination with L995F, exhibiting the same LD pattern as
133 N1570Y (Figure 1). These included two variants in codon 1874 (P1874S, P1874L), one
134 of which (P1874S) has previously been associated with pyrethroid resistance in the crop
135 pest *Plutella xylostella* [22]. The abundance of high-frequency non-synonymous variants
136 occurring in combination with L995F is striking for two reasons. First, *Vgsc* is a highly
137 conserved gene, expected to be under strong functional constraint and therefore purify-
138 ing selection, and so any non-synonymous variants are expected to be rare [11]. Second,
139 in contrast with L995F, we did not observe any high-frequency non-synonymous variants
140 occurring in combination with L995S. This contrast was highly significant when data on
141 all variants within the gene were considered: relative to haplotypes carrying the wild-type
142 L995 allele, the ratio of non-synonymous to synonymous nucleotide diversity (π_N/π_S) was
143 28.1 (95% CI [25.2, 31.2]) times higher among haplotypes carrying L995F but 1.5 (95%
144 CI [0.8, 2.2]) times higher among haplotypes carrying L995S. These results indicate that
145 L995F has substantially altered the selective regime for other amino acid positions within

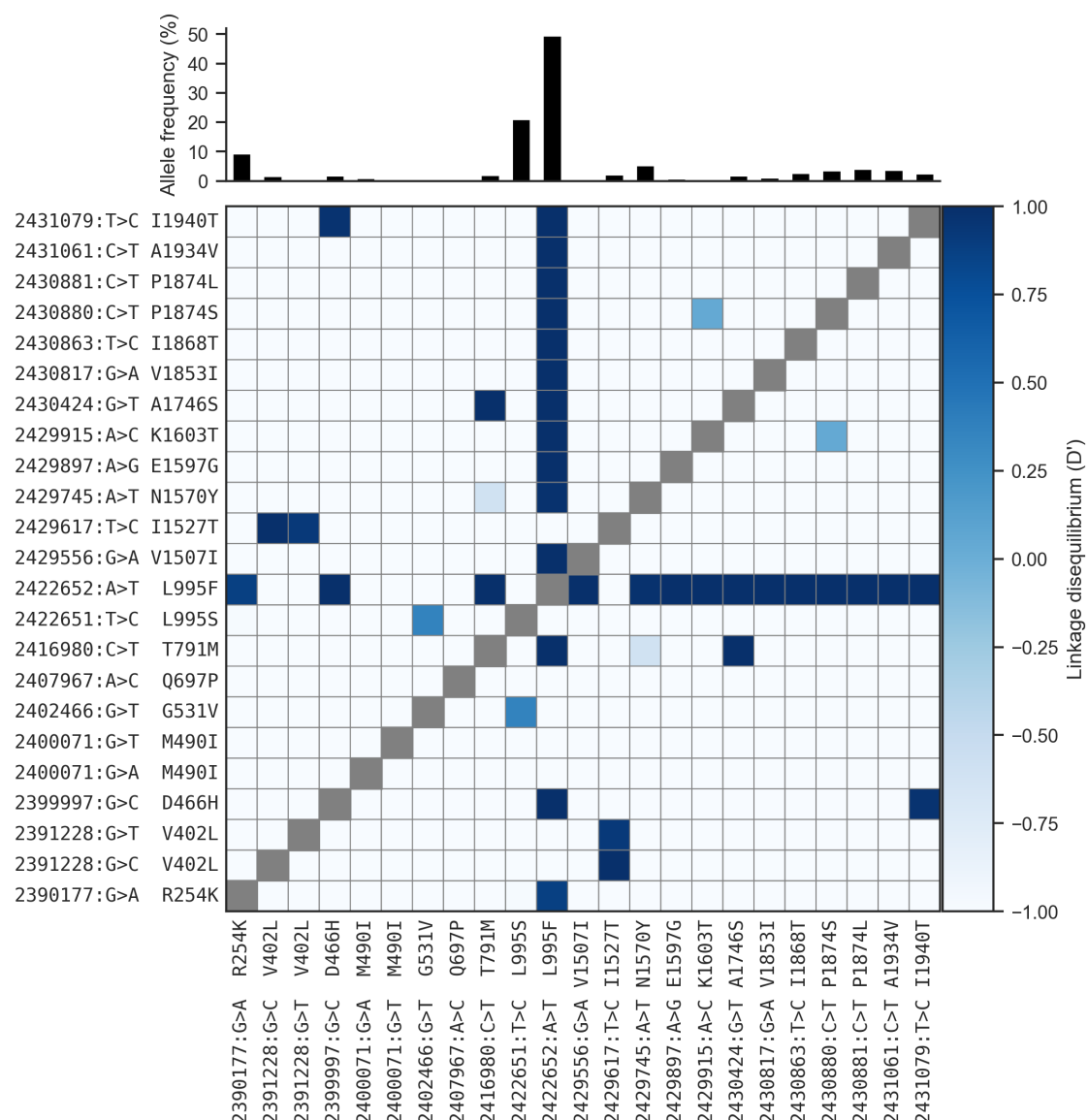


Figure 1. Linkage disequilibrium (D') between non-synonymous variants. A value of 1 indicates that two alleles are in perfect linkage, meaning that one of the alleles is only ever found in combination with the other. Conversely, a value of -1 indicates that two alleles are never found in combination with each other. The bar plot at the top shows the frequency of each allele within the Ag1000G phase 1 cohort. See Table 1 for population allele frequencies.

the protein. A number of secondary substitutions have occurred and risen in frequency, suggesting that they are providing some selective advantage in the presence of insecticide pressure.

A novel allele, I1527T, was present in *An. coluzzii* from Burkina Faso at 14% frequency. Codon 1527 occurs within trans-membrane segment IIIS6, immediately adjacent to residues within a predicted binding site for pyrethroid molecules, thus it is plausible that

152 I1527T could alter pyrethroid binding [23, 5]. We also found that the two variant alleles
153 affecting codon 402, both of which induce a V402L substitution, were in strong linkage
154 with I1527T ($D' \geq 0.8$; Figure 1), and almost all haplotypes carrying I1527T also carried a
155 V402L substitution. Substitutions in codon 402 have been found in a number of other insect
156 species and shown experimentally to confer pyrethroid resistance [5]. Because of the lim-
157 ited geographical distribution of these alleles, we hypothesize that the I1527T+V402L com-
158 bination represents a pyrethroid resistance allele that arose in West African *An. coluzzii*
159 populations. However, the L995F allele is at higher frequency (85%) in our Burkina Faso
160 *An. coluzzii* population, and is known to be increasing in frequency [24], therefore L995F
161 may provide a stronger resistance phenotype and is replacing I1527T+V402L.

162 The remaining 4 novel alleles (two separate nucleotide substitutions causing M490I;
163 A1125V; V1254I) did not occur in combination with any known resistance allele (Table 1).
164 All are private to a single population, and to our knowledge none have previously been
165 found in other species [12, 5].

166 Genetic backgrounds carrying resistance alleles

167 The Ag1000G data resource provides a rich source of information about the spread of
168 insecticide resistance alleles in any given gene, because data are available not only for
169 SNPs in protein coding regions, but also SNPs in introns and flanking intergenic regions,
170 and in neighbouring genes. These additional variants can be used to analyse the genetic
171 backgrounds (haplotypes) on which resistance alleles are found. In our initial report of
172 the Ag1000G phase 1 resource [20], we used 1710 biallelic SNPs from within the 73.5 kbp
173 *Vgsc* gene (1607 intronic, 103 exonic) to compute the number of SNP differences between
174 all pairs of 1530 haplotypes derived from 765 wild-caught mosquitoes. We then used
175 pairwise genetic distances to perform hierarchical clustering, and found that haplotypes
176 carrying resistance alleles in codon 995 were grouped into 10 distinct clusters, each with
177 near-identical haplotypes. Five of these clusters contained haplotypes carrying the L995F
178 allele (labelled F1-F5), and a further five clusters contained haplotypes carrying L995S
179 (labelled S1-S5).

180 To further investigate genetic backgrounds carrying resistance alleles, we used the
181 Ag1000G haplotype data to construct median-joining networks [25] (Figure 2). The net-

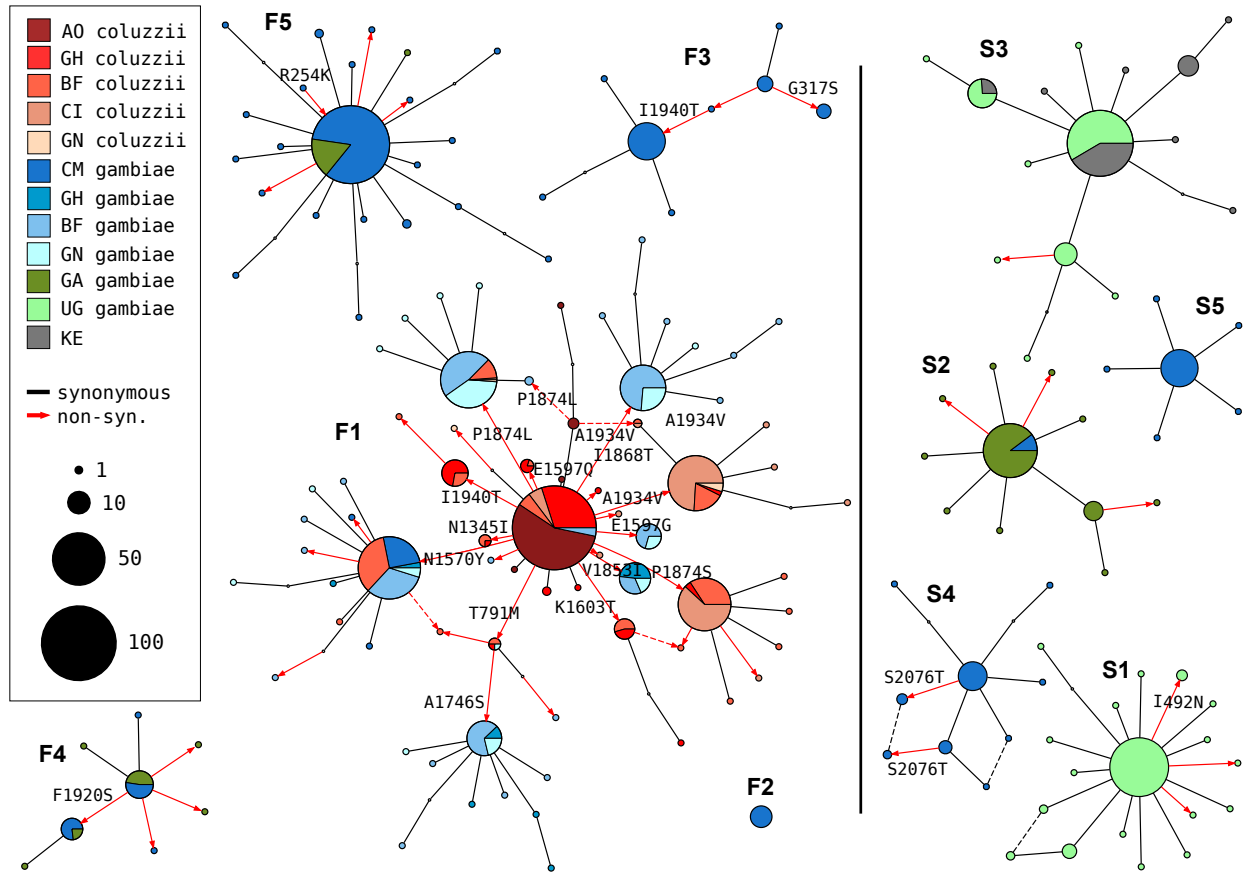


Figure 2. Haplotype networks. Median joining network for haplotypes carrying L995F (labelled F1-F5) or L995S variants (S1-S5) with a maximum edge distance of two SNPs. Labelling of network components is via concordance with hierarchical clusters discovered in [20]. Node size is relative to the number of haplotypes contained and node colour represents the proportion of haplotypes from mosquito populations/species - AO=Angola; BF=Burkina Faso; GN=Guinea; CM=Cameroon; GA=Gabon; UG=Uganda; KE=Kenya. Non-synonymous edges are highlighted in red and those leading to non-singleton nodes are labelled with the codon change, arrow head indicates direction of change away from the reference allele. Network components with fewer than three haplotypes are not shown.

work analysis improves on hierarchical clustering by allowing for the reconstruction and placement of intermediate haplotypes that may not be observed in the data. It also allows for non-hierarchical relationships between haplotypes, which may arise if recombination events have occurred between haplotypes. We constructed the network up to a maximum edge distance of 2 SNP differences, to ensure that each connected component captures a group of closely-related haplotypes. The resulting network contained 5 groups containing haplotypes carrying L995F, and a further 5 groups carrying L995S, in close correspondence with previous results from hierarchical clustering (96.8% overall concordance in assignment of haplotypes to groups).

191 The haplotype network brings into sharp relief the explosive radiation of amino acid sub-
 192 stitutions secondary to the L995F allele (Figure 2). Within the F1 group, nodes carrying
 193 non-synonymous variants radiate out from a central node carrying only L995F, suggest-
 194 ing that the central node represents the ancestral haplotype carrying L995F alone which
 195 initially came under selection, and these secondary variants have arisen subsequently as
 196 new mutations. Many of the nodes carrying secondary variants are large, consistent with
 197 positive selection and a functional role for these secondary variants as modifiers of the
 198 L995F resistance phenotype. The F1 network also allows us to infer multiple introgression
 199 events between the two species. The central (putatively ancestral) node contains hap-
 200 lotypes from individuals of both species, as do nodes carrying the N1570Y, P1874L and
 201 T791M variants. This structure is consistent with an initial introgression of the ancestral
 202 F1 haplotype, followed later by introgressions of haplotypes carrying secondary mutations.
 203 The haplotype network also illustrates the contrasting levels of non-synonymous varia-
 204 tion between L995F and L995S. Only two non-synonymous variants are present within the
 205 L995S groups, and both are at low frequency, thus may be neutral or mildly deleterious
 206 variants that are hitch-hiking on selective sweeps for the L995S allele.

207 The F1 group contained haplotypes from mosquitoes of both species, and from mosquitoes
 208 sampled in 4 different countries (Guinea, Burkina Faso, Cameroon, Angola) (Figure 3).
 209 The F4, F5 and S2 groups each contained haplotypes from both Cameroon and Gabon.
 210 The S3 group contained haplotypes from both Uganda and Kenya. The haplotypes
 211 within each of these groups were nearly identical across the entire span of the *Vgsc* gene
 212 ($\pi < 5.1 \times 10^{-5} bp^{-1}$). In contrast, diversity among wild-type haplotypes was two or-
 213 ders of magnitude greater (Cameroon *An. gambiae* $\pi = 1.4 \times 10^{-3} bp^{-1}$; Guinea-Bissau
 214 $\pi = 5.7 \times 10^{-3} bp^{-1}$). Thus it is reasonable to assume that each of these five groups con-
 215 tains descendants of an ancestral haplotype that carried a resistance allele and has risen in
 216 frequency due to selection for insecticide resistance. Given this assumption, these groups
 217 each provide evidence for adaptive gene flow between mosquito populations separated by
 218 considerable geographical distances.

219 A limitation of both the hierarchical clustering and network analyses is that they rely on
 220 genetic distances within a fixed genomic window from the start to the end of the *Vgsc* gene.
 221 *Anopheles* mosquitoes undergo homologous recombination during meiosis in both males

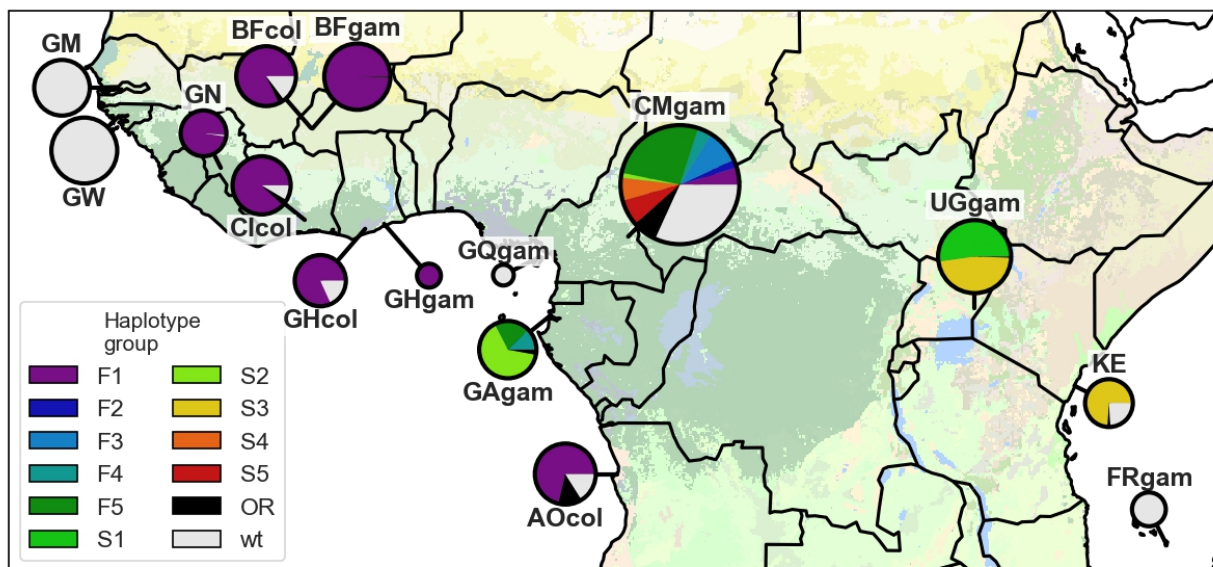


Figure 3. Map of haplotype frequencies. Each pie shows the frequency of different haplotype groups within one of the populations sampled. The size of the pie is proportional to the number of haplotypes sampled. The size of each wedge within the pie is proportional to the frequency of a haplotype group within the population. Haplotypes in groups F1-5 carry the L995F *kdr* allele. Haplotypes in groups S1-5 carry the L995S *kdr* allele. Haplotypes in group L1 carry the I1527T allele. Haplotypes in group L2 carry the M490I allele. Wild-type (*wt*) haplotypes do not carry any known or putative resistance alleles.

222 and females, and any recombination events that occurred within this genomic window
 223 could affect the way that haplotypes are grouped together in clusters or network compo-
 224 nents. In particular, recombination events could occur during the geographical spread of
 225 a resistance allele, altering the genetic background upstream and/or downstream of the
 226 allele itself. An analysis based on a fixed genomic window might then fail to infer gene flow
 227 between two mosquito populations, because haplotypes with and without a recombination
 228 event could be grouped separately, despite the fact that they share a recent common an-
 229 cestor. To investigate the possibility that recombination events may have affected our
 230 grouping of haplotypes carrying resistance alleles, we performed a windowed analysis of
 231 haplotype homozygosity, spanning *Vgsc* and up to a megabase upstream and downstream
 232 of the gene (Supplementary Figures S1, S2). This analysis supported a refinement of our
 233 initial grouping of haplotypes carrying resistance alleles. All haplotypes within groups S4
 234 and S5 were effectively identical on both the upstream and downstream flanks of the gene,
 235 but there was a region of divergence within the *Vgsc* gene itself that separated them in
 236 the fixed window analyses (Supplementary Figure S2). The 13.8 kbp region of divergence

occurred upstream of codon 995 and contained 8 SNPs that were fixed differences between S4 and S5. A possible explanation for this short region of divergence is that a gene conversion event has occurred within the gene, bringing a segment from a different genetic background onto the original genetic background on which the L995S resistance mutation occurred.

Positive selection for resistance alleles

To investigate evidence for positive selection on non-synonymous alleles, we performed an analysis of extended haplotype homozygosity (EHH) [26]. Haplotypes under recent positive selection will have increased rapidly in frequency, thus have had less time to be broken down by recombination, and should on average have longer regions of haplotype homozygosity relative to wild-type haplotypes. We defined a core region spanning *Vgsc* codon 995 and an additional 6 kbp of flanking sequence, which was the minimum required to differentiate the haplotype groups identified via clustering and network analyses. Within this core region, we found 18 distinct haplotypes at a frequency above 1% within the cohort. These included core haplotypes corresponding to each of the 10 haplotype groups carrying L995F or L995S alleles identified above, as well as a core haplotype carrying I1527T which we labelled L1 (due to it carrying the the wild-type leucine codon at position 995). We also found a core haplotype corresponding to a group of haplotypes from Kenya carrying an M490I allele, which we labelled as L2. All other core haplotypes we labelled as wild-type (*wt*). We then computed EHH decay for each core haplotype up to a megabase upstream and downstream of the core locus (Figure 4).

As expected, haplotypes carrying the L995F and L995S resistance alleles all experience a dramatically slower decay of EHH relative to wild-type haplotypes, supporting positive selection. Previous studies have found evidence for different rates of EHH decay between L995F and L995S haplotypes, suggesting differences in the timing and/or strength of selection [15]. However, we found no systematic difference in the length of shared haplotypes when comparing F1-5 (carrying L995F) against S1-5 (carrying L995S) (Supplementary Figure S3). There were, however, some differences between core haplotypes carrying the same allele. For example, shared haplotypes were significantly longer for S1 (median 1.091 cM, 95% CI [1.076 - 1.091]) versus other core haplotypes carrying L995S (e.g., S2 median

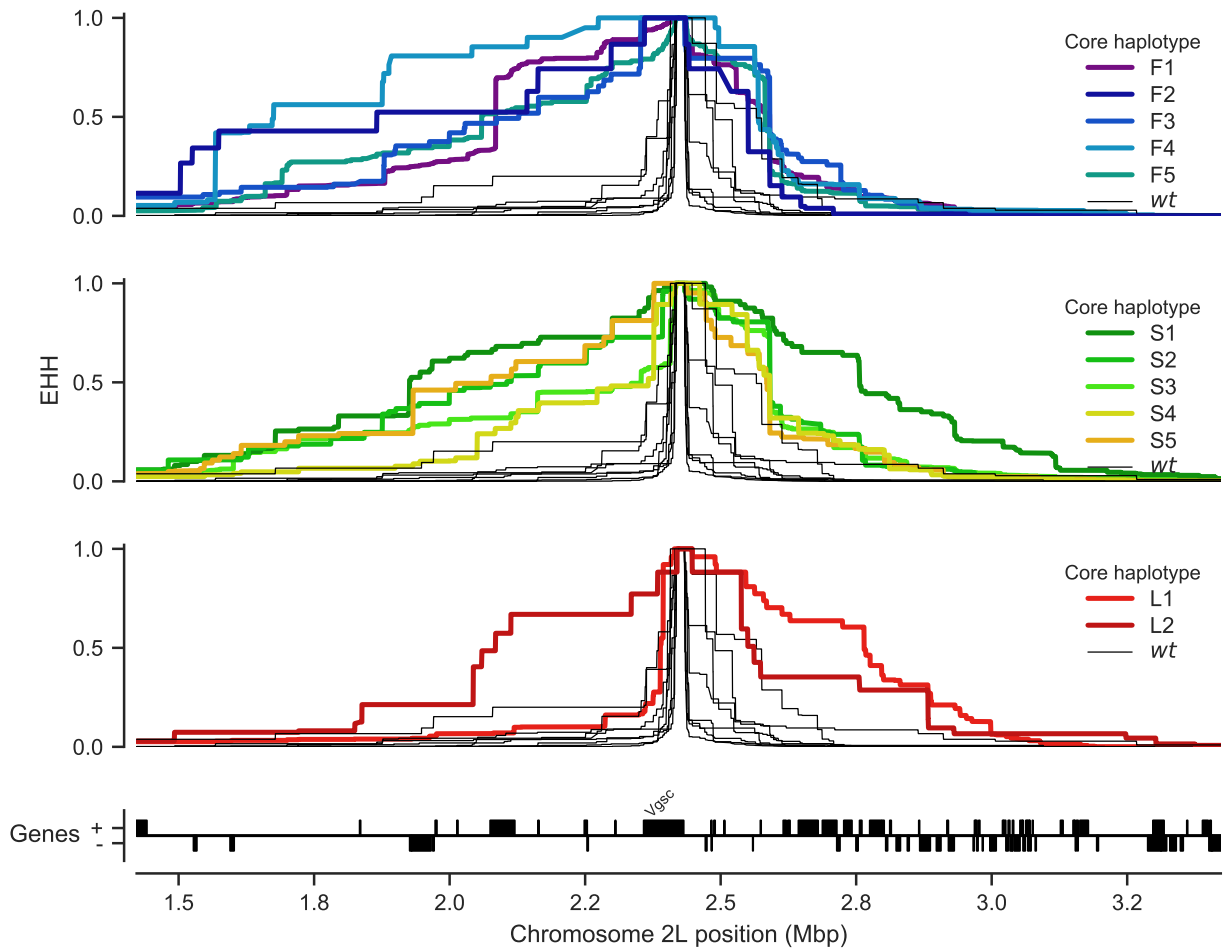


Figure 4. Evidence for positive selection on haplotypes carrying known or putative resistance alleles. Each panel plots the decay of extended haplotype homozygosity (EHH) for a set of core haplotypes centred on *Vgsc* codon 995. Core haplotypes F1-F5 carry the L995F allele; S1-S5 carry the L995S allele; L1 carries the I1527T allele; L2 carries the M490I allele. Wild-type (*wt*) haplotypes do not carry known or putative resistance alleles. A slower decay of EHH relative to wild-type haplotypes implies positive selection (each panel plots the same collection of wild-type haplotypes).

0.699 cM, 95% CI [0.696 - 0.705]; Supplementary Figure S3). Longer shared haplotypes indicate a more recent common ancestor, and thus some of these core haplotypes may have experienced more recent and/or more intense selection than others. The L1 haplotype carrying I1527T+V402L exhibited a slow decay of EHH on the downstream flank of the gene, similar to haplotypes carrying L995F and L995S, indicating that this combination of alleles has experienced positive selection. EHH decay on the upstream gene flank was faster, being similar to wild-type haplotypes, however there were two separate nucleotide substitutions encoding V402L within this group of haplotypes, and a faster EHH decay on this flank is consistent with recombination events bringing V402L alleles from differ-

ent genetic backgrounds together with an ancestral haplotype carrying I1527T. The L2 haplotype carrying M490I exhibited EHH decay on both flanks comparable to haplotypes carrying known resistance alleles. This could indicate evidence for selection on the M490I allele, however these haplotypes are derived from a Kenyan mosquito population where there is evidence for a severe recent bottleneck [20], and there were not enough wild-type haplotypes from Kenya with which to compare, thus this signal may also be due to the extreme demographic history of this population.

Discussion

Cross-resistance between pyrethroids and DDT

The VGSC protein is the physiological target of both pyrethroid insecticides and DDT [4]. The L995F and L995S alleles are known to increase resistance to both of these insecticide classes [6, 8]. By 2012, over half of African households owned at least one pyrethroid impregnated ITN and nearly two thirds of IRS programmes were using pyrethroids [2]. Pyrethroids were also introduced into agriculture in Africa prior to the scale-up of public health vector control programmes, and continue to be used on a variety of crops such as cotton [27]. DDT was used in Africa for several pilot IRS projects carried out during the first global campaign to eradicate malaria, during the 1950s and 1960s [11]. DDT is still approved for IRS use by WHO and remains in use in some locations, however within the last two decades pyrethroid use has been far more common and widespread. DDT was also used in agriculture from the 1940s, and although agricultural usage has greatly diminished since the 1970s, some usage remains [28]. In this study we reported evidence of positive selection on the L995F and L995S alleles, as well as the I1527T+V402L combination and possibly M490I. We also found 14 other non-synonymous substitutions that have arisen in association with L995F and appear to be positively selected. Given that pyrethroids have dominated public health insecticide use for two decades, it is reasonable to assume that the selection pressure on these alleles is primarily due to pyrethroids rather than DDT. It has previously been suggested that L995S may have been initially selected by DDT usage [15]. However, we did not find any systematic difference in the extent of haplotype homozygosity between these two alleles, suggesting that both alleles have been under selection over a

305 similar time frame. We did find some significant differences in haplotype homozygosity
 306 between different genetic backgrounds carrying resistance alleles, suggesting differences
 307 in the timing and/or strength of selection these may have experienced. However, there
 308 have been differences in the scale-up of pyrethroid-based interventions in different regions,
 309 and this could in turn generate heterogeneities in selection pressures. Nevertheless, it is
 310 possible that some if not all of the alleles we have reported provide some level of cross-
 311 resistance to DDT as well as pyrethroids, and we cannot exclude the possibility that
 312 earlier DDT usage may have contributed at least in part to their selection. The differing
 313 of resistance profiles to the two types of pyrethroids (type I, e.g., permethrin; and type
 314 II, e.g., deltamethrin) [29], will also affect the selection landscape. Further sampling and
 315 analysis is required to investigate the timing of different selection events and relate these
 316 to historical patterns of insecticide use in different regions.

317 **Resistance phenotypes for novel non-synonymous variants**

318 The sodium channel protein consists of four homologous domains (I-IV) each of which com-
 319 prises six transmembrane segments (S1-S6) connected by intracellular and extracellular
 320 loops [5]. Two pyrethroid binding sites have been predicted within the pore-forming mod-
 321 ules of the protein, the first (PyR1) involving residues from transmembrane segments IIS5
 322 and IIS6 and the internal linker between IIS4 and IIS5 (IIL45) [30], the second (PyR2)
 323 involving segments IS5, IS6, IIS6 and IL45 [23, 5]. Many of the amino acid substitutions
 324 known to cause pyrethroid resistance in insects affect residues within one of these two
 325 pyrethroid binding sites, and thus can directly alter pyrethroid binding [5]. For example,
 326 the L995F and L995S substitutions occur in segment IIS6 and belong to binding site PyR2
 327 [23]. The I1527T substitution that we discovered in *An. coluzzii* mosquitoes from Burk-
 328 ina Faso occurs in segment IIS6 and is immediately adjacent to two pyrethroid-sensing
 329 residues in site PyR1 [5]. It is thus plausible that pyrethroid binding could be altered by
 330 this substitution. The I1527T substitution (*M. domestica* codon 1532) has been found in
 331 *Aedes albopictus* [31], and substitutions in the nearby codon 1529 (*M. domestica* codon
 332 1534) have been reported in *Aedes albopictus* and in *Aedes aegypti* where it was found to be
 333 associated with pyrethroid resistance [5, 32, 33]. We found the I1527T allele in tight link-
 334 age with two alleles causing a V402L substitution (*M. domestica* codon 410). Substitutions

335 in codon 402 have been found in multiple insect species and are by themselves sufficient to
336 confer pyrethroid resistance [5]. Codon 402 is within segment IS6, immediately adjacent
337 to a pyrethroid sensing residue in site PyR2. The fact that we find I1527T and V402L in
338 such tight mutual association is intriguing because (a) these two residues appear to affect
339 different pyrethroid binding sites, and (b) haplotypes carrying V402L alone should also
340 have been positively selected and thus be present in one or more populations.

341 A number of substitutions in segments of the protein that are not involved in either
342 of the two pyrethroid binding sites have also been shown to confer pyrethroid resistance.
343 For example, the N1570Y substitution causes substantially enhanced pyrethroid resistance
344 when combined with L995F, although codon 1570 occurs in the internal linker between
345 domains III and IV (LIII/IV) [23]. Computer modelling of the protein structure has sug-
346 gested that substitutions in codon 1570 could allosterically alter site PyR2 and thus affect
347 pyrethroid binding [23]. In addition to N1570Y, we found thirteen other substitutions at
348 appreciable frequency occurring almost exclusively in association with L995F (Table 1;
349 Figure 1). Of these, two (D466H, E1597G) occurred in the larger internal linkers between
350 protein domains, one (R254K) occurred within a smaller internal linker between domain
351 subunits, two (T791M, K1603T) occurred within an outer (“voltage-sensing”) transmem-
352 brane segment, one (A1746S) occurred within an inner (“pore-forming”) transmembrane
353 segment, and the remaining seven occurred in the internal carboxyl-terminal tail. Thus
354 there is no simple pattern regarding where these variants occur within the protein struc-
355 ture. Further work is required to confirm which of these substitutions affect pyrethroid
356 resistance, and to determine whether they allosterically modify a pyrethroid binding site
357 in a similar vein to N1570Y, or whether they provide some other benefit such as compen-
358 sating for a deleterious effect of L995F on normal nervous system function. The novel
359 M490I substitution, found on the Kenyan L2 haplotypic background potentially under se-
360 lection, also occurs in an internal linker between protein domains (LI/II). However, M490I
361 did not occur in association with L995F or any other non-synonymous substitutions. It is
362 plausible that substitutions outside of pyrethroid binding sites could independently confer
363 an insecticide resistance phenotype, because there are several known examples in other
364 insect species [5]. Work in other species has also suggested that pyrethroid resistance sub-
365 stitutions could act not by altering pyrethroid binding but by altering the channel gating

kinetics or the voltage-dependence of activation [5]. Thus there are a number of potential mechanisms by which a pyrethroid resistance phenotype can be obtained, and clearly much remains to be unravelled regarding the molecular biology of pyrethroid resistance in this gene.

Design of genetic assays for surveillance of pyrethroid resistance

Entomological surveillance teams in Africa regularly genotype mosquitoes for resistance alleles in *Vgsc* codon 995, and use those results as an indicator for the presence of pyrethroid resistance alongside results from insecticide resistance bioassays. They typically do not, however, sequence the gene or genotype any other polymorphisms within the gene. Thus if there are other polymorphisms within the gene that cause or significantly enhance pyrethroid resistance, these will not be detected. Also, if a codon 995 resistance allele is observed, there is no way to know whether the allele is on a genetic background that has also been observed in other mosquito populations, and thus no way to investigate whether resistance alleles are emerging locally or being imported from elsewhere. Whole-genome sequencing of individual mosquitoes clearly provides data of sufficient resolution to answer these questions, and could be used to provide ongoing resistance surveillance. The cost of whole-genome sequencing continues to fall, with the present cost being approximately 50 GBP to obtain ~30× coverage of an individual *Anopheles* mosquito genome with 150 bp paired-end reads. However, to achieve substantial spatial and temporal coverage of mosquito populations, it is currently cheaper and more practical to develop targeted genetic assays for resistance outbreak surveillance. Technologies such as amplicon sequencing [34] could scale to tens of thousands of mosquitoes at low cost and could be implemented using existing platforms in national molecular biology facilities.

To facilitate the development of targeted genetic assays for surveillance of *Vgsc*-mediated pyrethroid resistance, we have produced several supplementary data tables. In Supplementary Table 1 we list all 64 non-synonymous variants found within the *Vgsc* gene in this study, with population allele frequencies. In Supplementary Table 2 we list 771 biallelic SNPs, within the *Vgsc* gene and up to 10 kbp upstream or downstream, that are potentially informative regarding which haplotype group a resistance haplotype belongs to, and thus could be used for tracking the spread of resistance. This table includes the allele

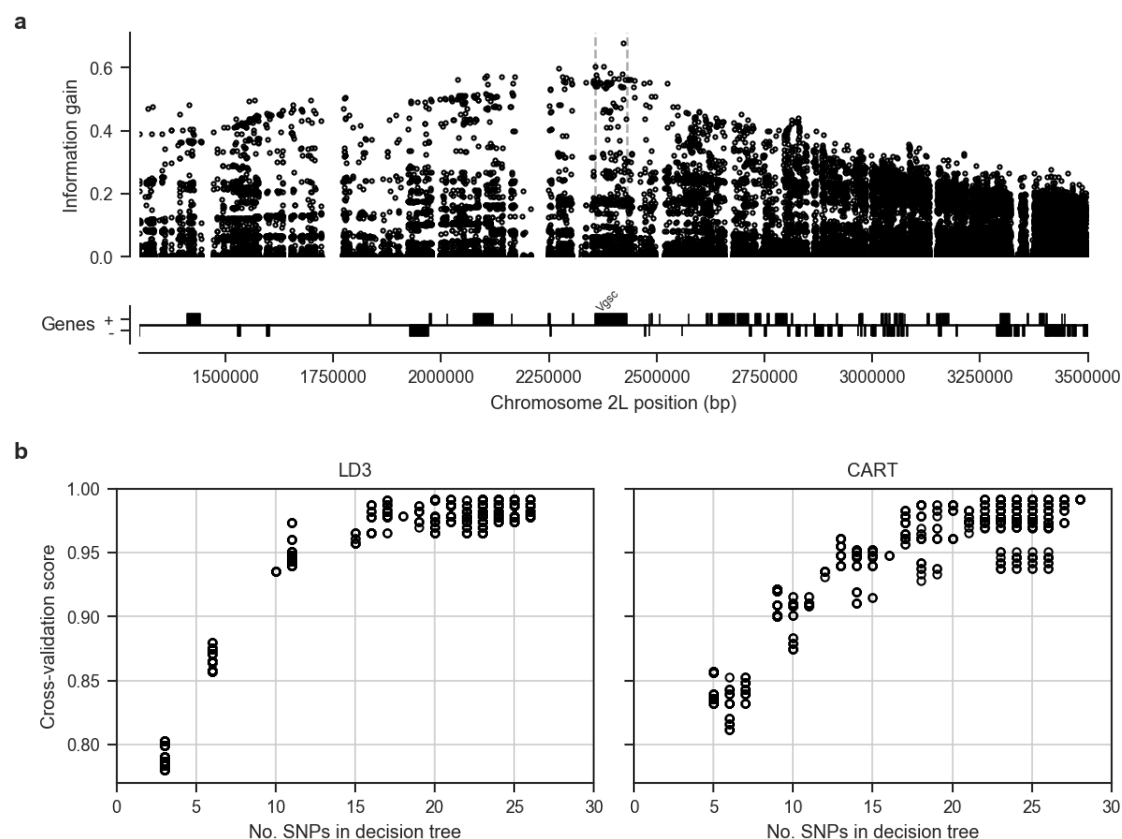


Figure 5. Informative SNPs for haplotype surveillance. **a**, Each data point represents a single SNP. The information gain value for each SNP provides an indication of how informative the SNP is likely to be if used as part of a genetic assay for testing whether a mosquito carries a resistance haplotype, and if so, which haplotype group it belongs to. **b**, Number of SNPs required to accurately predict which group a resistance haplotype belongs to. Each data point represents a single decision tree. Decision trees were constructed using either the LD3 (left) or CART (right) algorithm for comparison. Accuracy was evaluated using 10-fold stratified cross-validation.

frequency within each of the 12 haplotype groups defined here, to aid in identifying SNPs that are highly differentiated between two or more haplotype groups. We also provide Supplementary Table 3 which lists all 8,297 SNPs found within the *Vgsc* gene and up to 10 kbp upstream or downstream, which might need to be taken into account as flanking variation when searching for PCR primers to amplify a SNP of interest. To provide some indication for how many SNPs would need to be assayed in order to track the spread of resistance, we used haplotype data from this study to construct decision trees that could classify which of the 12 groups a given haplotype belongs to (Figure 5). This analysis suggested that it should be possible to construct a decision tree able to classify haplotypes with >95% accuracy by using 20 SNPs or less. In practice, more SNPs would be needed, to provide some redundancy, and also to type non-synonymous polymorphisms in

407 addition to identifying the genetic background. However, it is still likely to be well within
408 the number of SNPs that could be assayed in a single multiplex via amplicon sequencing.
409 Thus it should be feasible to produce low-cost, high-throughput genetic assays for tracking
410 the spread of pyrethroid resistance. If combined with a limited amount of whole-genome
411 sequencing at sentinel sites, this should also allow the identification of newly emerging
412 resistance outbreaks.

413 **Methods**

414 **Code**

415 All scripts and Jupyter Notebooks used to generate analyses, figures and tables are avail-
416 able from the GitHub repository <https://github.com/malariagen/agam-vgsc-report>.

417 **Data**

418 We used variant calls from the Ag1000G Phase 1 AR3 data release ([https://www.malariagen.](https://www.malariagen.net/data/ag1000g-phase1-ar3)
419 [net/data/ag1000g-phase1-ar3](https://www.malariagen.net/data/ag1000g-phase1-ar3)) and phased haplotype data from the Ag1000G Phase 1
420 AR3.1 data release (<https://www.malariagen.net/data/ag1000g-phase1-ar3.1>). Vari-
421 ant calls from Ag1000G Phase 1 are also available from the European Nucleotide Archive
422 (ENA; <http://www.ebi.ac.uk/ena>) under study PRJEB18691.

423 **Data collection and processing**

424 For detailed information on Ag1000G WGS sample collection, sequencing, variant calling,
425 quality control and phasing, see [20]. In brief, *An. gambiae* and *An. coluzzii* mosquitoes
426 were collected from eight countries across Sub-Saharan Africa: Angola, Burkina Faso,
427 Cameroon, Gabon, Guinea, Guinea Bissau, Kenya and Uganda. From Angola just *An.*
428 *coluzzii* were sampled, Burkina Faso had samples of both *An. gambiae* and *An. coluzzii*
429 and all other populations consisted of purely *An. gambiae*, except for Kenya and Guinea
430 Bissau where species status is uncertain [20]. Mosquitoes were individually whole genome
431 sequenced on the Illumina HiSeq 2000 platform, generating 100bp paired-end reads. Se-
432 quence reads were aligned to the *An. gambiae* AgamP3 reference genome assembly [35].
433 Aligned bam files underwent improvement, before variants were called using GATK Uni-

434 fiedGenotyper. Quality control included removal of samples with mean coverage $\leq 14\times$
435 and filtering of variants with attributes that were correlated with Mendelian error in ge-
436 netic crosses.

437 The Ag1000G variant data was functionally annotated using the SnpEff v4.1b soft-
438 ware [36]. Non-synonymous *Vgsc* variants were identified as all variants in transcript
439 AGAP004707-RA with a SnpEff annotation of “missense”. The *Vgsc* gene is known to
440 exhibit alternative splicing [4], however at the time of writing the *An. gambiae* gene an-
441 notations did not include the alternative transcripts reported by Davies et al. We wrote
442 a Python script to check for the presence of variants that are synonymous according to
443 transcript AGAP004707-RA but non-synonymous according to one of the other transcripts
444 present in the gene annotations or in the set reported by Davies et al. Supplementary Ta-
445 ble 1 includes the predicted effect for all SNPs that are non-synonymous in one or more
446 of these transcripts. None of the variants that are non-synonymous in a transcript other
447 than AGAP004707-RA were found to be above 5% frequency in any population.

448 For ease of comparison with previous work on *Vgsc*, pan Insecta, in Table 1 and Supple-
449 mentary Table 1 we report codon numbering for both *An. gambiae* and *Musca domestica*
450 (the species in which the gene was first discovered). The *M. domestica* *Vgsc* sequence
451 (EMBL accession X96668 [9]) was aligned with the *An. gambiae* AGAP004707-RA se-
452 quence (AgamP4.4 gene-set) using the Mega v7 software package [37]. A map of equiva-
453 lent codon numbers between the two species for the entire gene can be download from the
454 MalariaGEN website ([https://www.malariagen.net/sites/default/files/content/](https://www.malariagen.net/sites/default/files/content/blogs/domestica_gambiae_map.txt)
455 [blogs/domestica_gambiae_map.txt](https://www.malariagen.net/sites/default/files/content/blogs/domestica_gambiae_map.txt)).

456 Haplotypes for each chromosome of each sample were estimated (phased) using using
457 phase informative reads (PIRs) and SHAPEIT2 v2.r837 [38], see [20] supplementary text
458 for more details. The SHAPEIT2 algorithm is unable to phase multi-allelic positions,
459 therefore the two multi-allelic non-synonymous SNPs within the *Vgsc* gene, altering codons
460 V402 and M490, were phased onto the biallelic haplotype scaffold using MVNcall v1.0 [39].
461 Conservative filtering applied to the genome-wide callset had removed one of the three
462 known insecticide resistance conferring *kdr* variants, N1570Y [10]. Manual inspection of
463 the read alignment revealed that the SNP call could be confidently made, and it was
464 added back into the data set and then also phased onto the haplotypes using MVNcall.

Lewontin's D' [40] was used to compute the linkage disequilibrium (LD) between all pairs of non-synonymous *Vgsc* mutations.

Haplotype networks

Haplotype networks were constructed using the median-joining algorithm [25] as implemented in a Python module available from <https://github.com/malariagen/agam-vgsc-report>. Haplotypes carrying either L995F or L995S mutations were analysed with a maximum edge distance of two SNPs. Networks were rendered with the Graphviz library and a composite figure constructed using Inkscape. Non-synonymous edges were highlighted using the SnpEff annotations [36].

Positive selection

Core haplotypes were defined on a 6,078 bp region spanning *Vgsc* codon 995, from chromosome arm 2L position 2,420,443 and ending at position 2,426,521. This region was chosen as it was the smallest region sufficient to differentiate between the ten genetic backgrounds carrying either of the known resistance alleles L995F or L995S. Extended haplotype homozygosity (EHH) was computed for all core haplotypes as described in [26] using scikit-allel version 1.1.9 [41], excluding non-synonymous and singleton SNPs. Analyses of haplotype homozygosity in moving windows (Supplementary Figs. S1, S2) and pairwise haplotype sharing (Supplementary Figure S3) were performed using custom Python code available from <https://github.com/malariagen/agam-vgsc-report>.

Design of genetic assays for surveillance of pyrethroid resistance

To explore the feasibility of identifying a small subset of SNPs that would be sufficient to identify each of the genetic backgrounds carrying known or putative resistance alleles, we started with an input data set of all SNPs within the *Vgsc* gene or in the flanking regions 20 kbp upstream and downstream of the gene. Each of the 1530 haplotypes in the Ag1000G Phase 1 cohort was labelled according to which core haplotype it carried, combining all core haplotypes not carrying known or putative resistance alleles together as a single "wild-type" group. Decision tree classifiers were then constructed using scikit-learn version 0.19.0 [42] for a range of maximum depths, repeating the tree construction process

10 times for each maximum depth with a different initial random state. The classification accuracy of each tree was evaluated using stratified 5-fold cross-validation.

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613 **Supplementary figures**

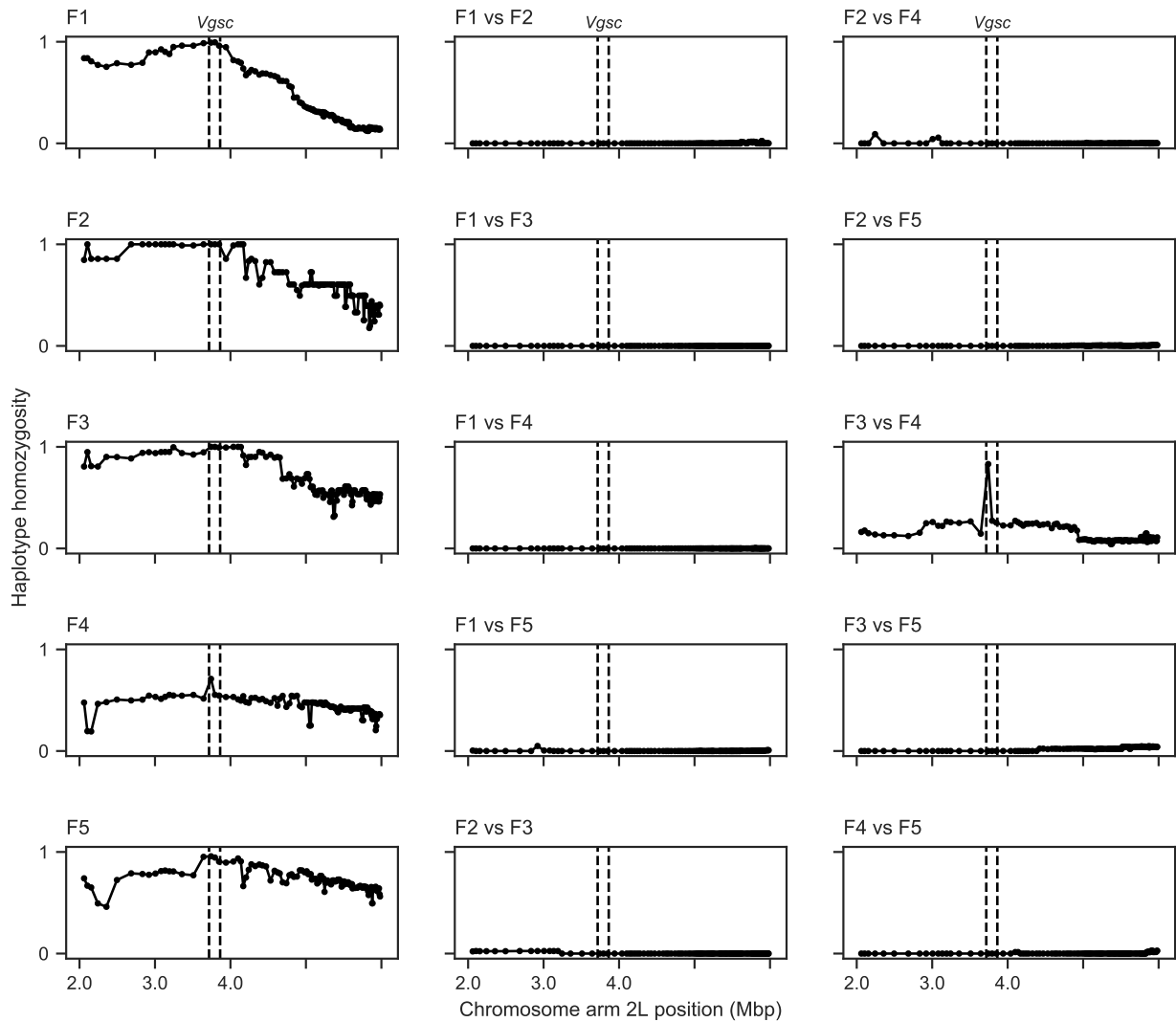


Figure S1. Windowed analysis of haplotype homozygosity for genetic backgrounds carrying the L995F allele. Each sub-plot shows the fraction of haplotype pairs that are identical within half-overlapping moving windows of 1000 SNPs. Each sub-plot in the left-hand column shows homozygosity for haplotype pairs within one of the haplotype groups identified by the network analysis. Sub-plots in the central and right-hand columns show homozygosity for haplotype pairs between two haplotype groups. If two haplotype groups are truly unrelated, haplotype homozygosity between them should be close to zero across the whole genome region. Dashed vertical lines show the location of the *Vgsc* gene.

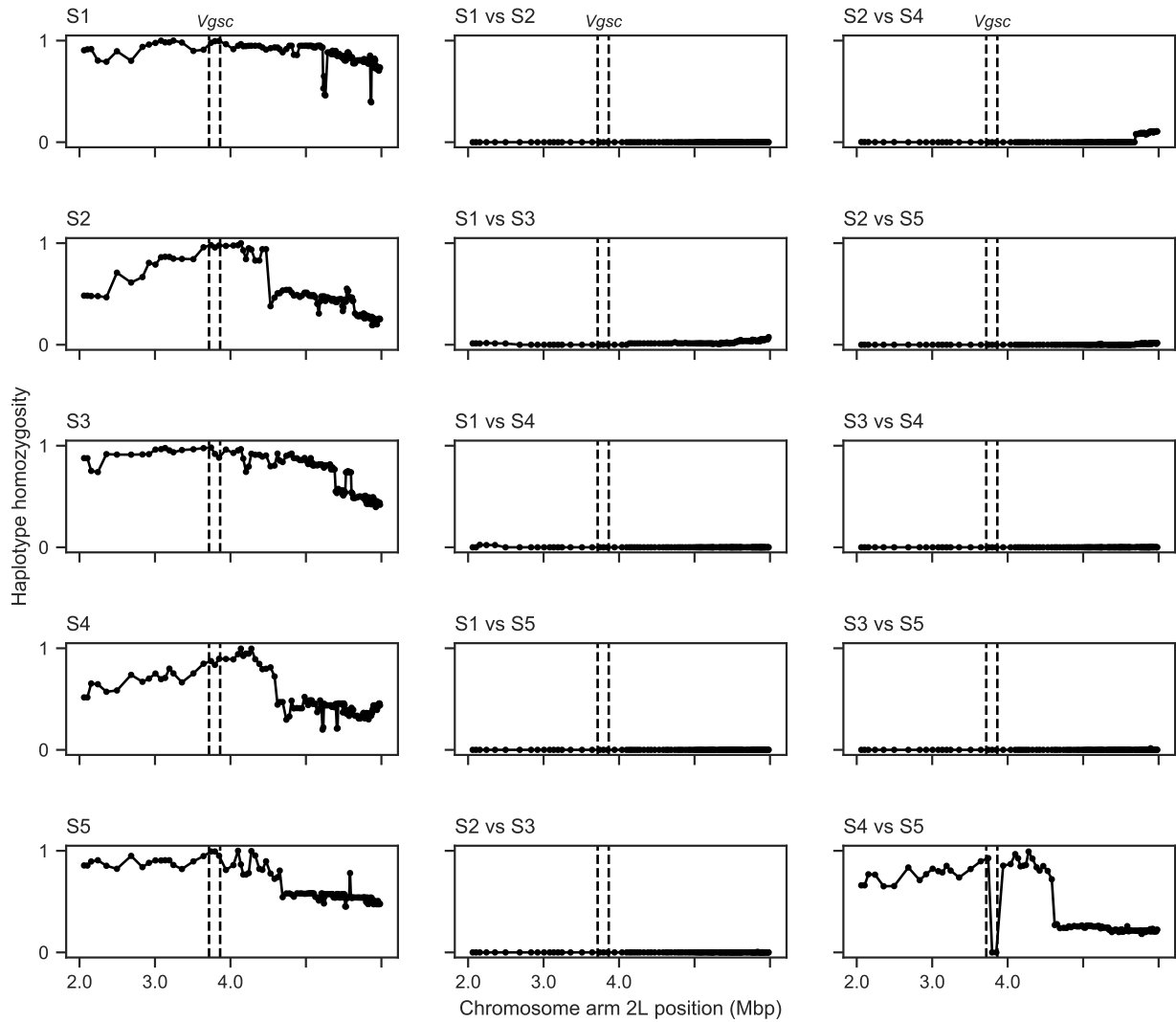


Figure S2. Windowed analysis of haplotype homozygosity for genetic backgrounds carrying the L995S allele. See Supplementary Figure S1 for explanation. Haplotype homozygosity is high between groups S4 and S5 on both flanks of the gene, indicating that haplotypes from both groups are in fact closely related.

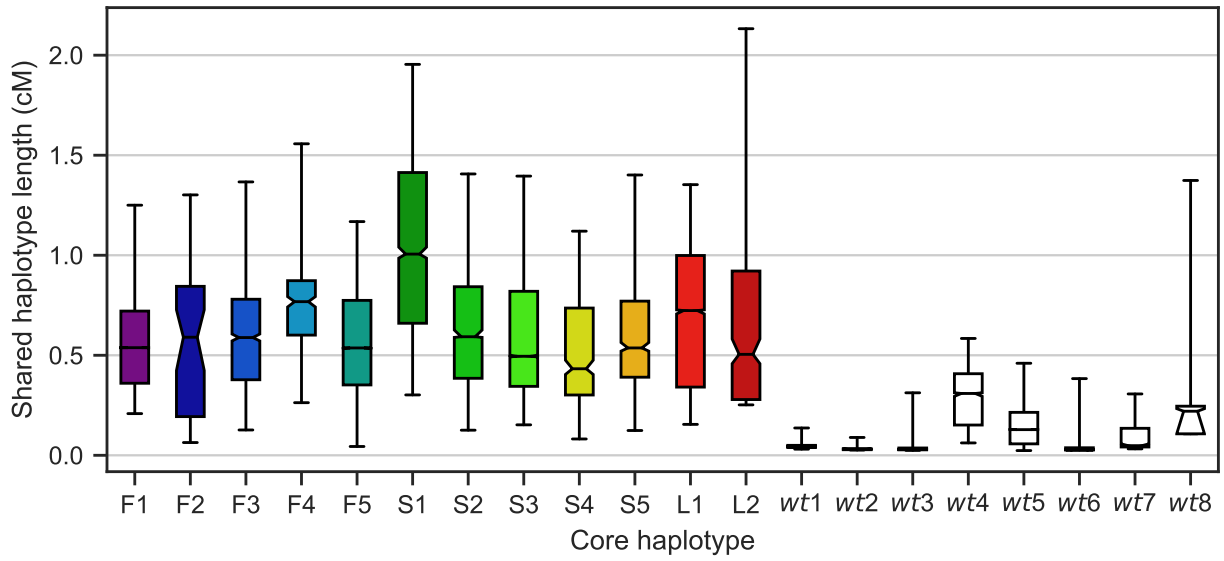


Figure S3. Shared haplotype length. Each bar shows the distribution of shared haplotype lengths between all pairs of haplotypes with the same core haplotype. For each pair of haplotypes, the shared haplotype length is computed as the region extending upstream and downstream from the core locus (*Vgsc* codon 995) over which haplotypes are identical at all non-singleton variants. The *Vgsc* gene sits on the border of pericentromeric heterochromatin and euchromatin, and we assume different recombination rates in upstream and downstream regions. The shared haplotype length is expressed in centiMorgans (cM) assuming a constant recombination rate of 2.0 cM/Mb on the downstream (euchromatin) flank and 0.6 cM/Mb on the upstream (heterochromatin) flank. Bars show the inter-quartile range, fliers show the 5-95th percentiles, horizontal black line shows the median, notch in bar shows the 95% bootstrap confidence interval for the median. Haplotypes F1-5 each carry the L995F resistance allele. Haplotypes S1-5 each carry the L995S resistance allele. Haplotype L1 carries the I1527T allele. Haplotype L2 carries the M490I allele. Wild-type (*wt*) haplotypes do not carry any known or putative resistance alleles.